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(21) International Application Number: PCT/EP99/04723 (22) International Filing Date: 5 July 1999 (05.07.99) (30) Priority Data: 09/113,699 10 July 1998 (10.07.98) US (71) Applicant (for all designated States except US): BOX O3 INTERNATIONAL [CH/CH]; Hintere Dorfasse 9, CH-3073 Gümligen (CH). (72) Inventor; and (75) Inventor/Applicant (for US only): DUROSELLE, Patrick [FR/FR]; 18 Rue Nungesser et Coli, F-69008 Lyon (FR). (74) Agent: BOVARD LTD.; Optingenstrasse 16, CH-3000 Bern 25 (CH).		(81) Designated States: AE, AL, AM, AT, AT (Utility model), AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, CZ (Utility model), DE, DE (Utility model), DK, DK (Utility model), EE, EE (Utility model), ES, FI, FI (Utility model), GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SK (Utility model), SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZA, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SL, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG). Published <i>With international search report.</i>
(54) Title: A METHOD FOR DISINFECTING AND STERILIZING MICROBIAL CONTAMINATED MATERIALS (57) Abstract <p>The method for disinfecting and sterilizing microbial contaminated or infectious materials, e.g. medical instruments, waste from medical laboratories or hospitals, comprises the following steps: (a) loading the material for disinfecting and sterilizing into a vacuum-proof sterilization chamber; (b) introducing liquid hydrogen peroxide into the sterilization chamber so as to penetrate the material; (c) introducing liquid acetic acid into the sterilization chamber so as to penetrate the material; (d) evacuating gas from the sterilization chamber so as to evaporate the liquid at least partially; (e) introducing gaseous ozone into the sterilization chamber; (f) treating the material in a sterilization chamber for a sufficient time period so as to disinfect and sterilize the materials. The germicide potency of the method is markedly improved in comparison with the method where only compound O₃ is used alone or in combination with the germicide H₂O₂.</p>		

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A Method for Disinfecting and Sterilizing Microbial Contaminated Materials

The present invention relates to a method for disinfecting and sterilizing microbial contaminated materials, e.g. medical devices, instruments, waste from laboratories or hospitals comprising blood or blood components or other biological material under use of a combination of *in situ* generated peracetic acid and ozone.

For disinfecting of medical devices, working tools or infectious waste material in the medical field, several methods using ozone are suggested according to prior art. The main problem of known methods has been the long period for treating the material containing pathogenic agents such as bacteria and viruses. With a shorter treatment problems occurred with material containing biological liquids such as blood or components thereof (probably due to hemoglobin presence) because an amount of resistant spores survived. It was found that an improvement can be attained if a combination of ozone and peracetic acid was used in a humidified gaseous phase. Up to now peracetic acid has been used in liquid form for decontaminating medical wastes (US Patent 5,374,394) and the combination of peracetic acid and ozone as a disinfectant has never been used in liquid form. Another approach is described in the document JP-7-136236 wherein hot vapor is injected into a sterilization chamber. The use of a hot oxidizing agent can be irritating and dangerous in the case of a defect in the chamber.

The present invention is related to a method for disinfecting and sterilizing microbial contaminated or infectious materials, e.g. medical instruments, waste from medical laboratories or hospitals. The method comprises the steps:

- (a) loading the material for disinfecting and sterilizing into a pressure-proof sterilization chamber
- (b) introducing liquid hydrogen peroxide into the sterilization chamber so as to penetrate the material;
- (c) introducing liquid acetic acid into the sterilization chamber so as to penetrate the material;

- (d) evacuating gas from the sterilization chamber so as to evaporate the liquid at least partially;
 - (e) introducing gaseous ozone into the sterilization chamber;
 - (f) treating the material in the sterilization chamber for a sufficient time period so
- 5 as to disinfect and sterilize the materials

The method can be carried out in treatment devices as described in the patent documents EP-A- 0 664 715 and EP-A-0 761 237, which can be adapted to the method according to present invention.

The method of the present invention is carried out, as a rule, at a

10 temperature of about 15 °C to 35 °C and preferably at ambient temperature.

The hydrogen peroxide and the acetic acid can be introduced enclosed in two separate ampoules having such properties that after the evacuation of the chamber they are burst with release of the hydrogen peroxide and the acetic acid.

15 Alternatively the hydrogen peroxide and the acetic acid can be introduced enclosed in a container having at least two compartments being constructed in such a manner that the two compartments release their contents during the evacuation of the sterilizing chamber.

The method of the invention can be carried out in two or more cycles

20 by feeding at least two times hydrogen peroxide, the acetic acid and/or ozone into the sterilization chamber.

Alternatively the sterilization chamber can be equipped with storage and feeding means for feeding continuously hydrogen peroxide, acetic acid and ozone in several process cycles.

25 The molar ratio of acetic acid to ozone according to the invention is as a rule about 3/1 to 1/3 and the humidity in the chamber is at least 10 %.

The infectious waste material treated according to the method of the present invention can be solid material or a mixture of solid and liquid material. For disinfecting waste material safely, it is ground or broken up in other wise before, during or after the treatment with the disinfecting agents (hydrogen
5 peroxide and acetic acid).

Preferably a stable, storable and shippable disinfecting and sterilizing combination of agents is used together with ozone in the method of the invention. The combination possesses improved properties and comprises a two-part system: the first part consists of a mixture of acetic acid and water, and
10 the second consists of hydrogen peroxide and water. In this solution the peracetic acid is formed *in situ*. With this measure it is possible to take advantage of the outstanding disinfecting and sterilizing properties of peracetic acid without exposure to the danger of explosion and to the strong irritating properties to the skin and the eyes. The application of the peracetic acid
15 forming combination with ozone is carried out in a gaseous phase under reduced pressure (pressure < atmospheric pressure) and at ambient temperature. Hydrogen peroxide reacts with acetic acid to form of peracetic acid and water:



20 This is an equilibrium which can be shifted if the reaction takes place in the presence of a catalytic amount of acid (e.g. sulfuric acid). After termination of the disinfecting process the remaining ozone is destroyed, and, if necessary, the acid can be neutralized.

The present invention is illustrated with sterilizing tests carried out in
25 a sterilizing apparatus of the type "BOX O3" of Carbagas Aktiengesellschaft CH 3097 Liebefeld (Switzerland). This apparatus was designed for disinfecting infectious waste material. Normally in this device the waste material is ground , evacuated and treated one or several times with gaseous ozone in a sterilizing chamber. According to the present invention *in situ*-formed peracetic acid is
30 used additionally. For comparison, instead of peracetic acid only hydrogen

peroxide was used. For the test, pads loaded with infected sheep's blood were used. The results are listed in the table below.

Table: Number of viable and sporular bacterial forms destroyed in a sterilization chamber on a pad in the presence of blood of a sheep (100 µl per germ-carrier)

micro-organism	Number of cycles	Formation time	Compound 1	Compound 2	Reduction of micro-organisms
<i>S. aureus</i> +100µl of sheep's blood	3	900 s	H ₂ O ₂	-	3.87 log ₁₀
<i>S. aureus</i> +100µl of sheep's blood	3	900 s	H ₂ O ₂	CH ₃ COOH	7.90 log ₁₀
<i>S. aureus</i> +100µl of sheep's blood	3	900 s	H ₂ O ₂	CH ₃ COOH	<8.11 log ₁₀
<i>E. hirae</i> +100µl of sheep's blood	3	900 s	H ₂ O ₂	-	2.23 log ₁₀
<i>E. hirae</i> +100µl of sheep's blood	4	600 s	H ₂ O ₂	CH ₃ COOH	≥7 log ₁₀
<i>E. hirae</i> +100µl of sheep's blood	3	900 s	H ₂ O ₂	CH ₃ COOH	<7.8 log ₁₀
<i>E. coli</i> +100µl of sheep's blood	3	900 s	H ₂ O ₂	CH ₃ COOH	<7.69 log ₁₀
<i>E. coli</i> +100µl of sheep's blood	5	480 s	H ₂ O ₂	-	1.38 log ₁₀
<i>E. coli</i> +100µl of sheep's blood	3	900 s	H ₂ O ₂	CH ₃ COOH	≥8 log ₁₀
<i>E. coli</i> +100µl of sheep's blood	3	900 s	H ₂ O ₂	CH ₃ COOH	<8 log ₁₀

<i>P. aeruginosa</i> +100µl of sheep's blood	5	480 s	H ₂ O ₂	-	1.47 log ₁₀
<i>P. aeruginosa</i> +100µl of sheep's blood	4	900 s	H ₂ O ₂	CH ₃ COOH	7 log ₁₀
<i>P. aeruginosa</i> +100µl of sheep's blood	3	900 s	H ₂ O ₂	CH ₃ COOH	≥8.25 log ₁₀
<i>P. aeruginosa</i> +100µl of sheep's blood	3	900 s	H ₂ O ₂	CH ₃ COOH	8.47 log ₁₀
<i>M. smegmatis</i> +100µl of sheep's blood	3	900 s	H ₂ O ₂	-	2.47 log ₁₀
<i>M. smegmatis</i> +100µl of sheep's blood	4	600 s	H ₂ O ₂	CH ₃ COOH	≤7 log ₁₀
<i>M. smegmatis</i> +100µl of sheep's blood	3	900 s	H ₂ O ₂	CH ₃ COOH	≤7.6 log ₁₀
<i>M. smegmatis</i> +100µl of sheep's blood	3	900 s	H ₂ O ₂	CH ₃ COOH	≤8 log ₁₀
<i>C. albicans</i> +100µl of sheep's blood	3	900 s	H ₂ O ₂	-	1.8 log ₁₀
<i>C. albicans</i> +100µl of sheep's blood	3	900 s	H ₂ O ₂	CH ₃ COOH	≥8.04 log ₁₀
<i>C. albicans</i> +100µl of sheep's blood	3	900 s	H ₂ O ₂	CH ₃ COOH	≥8.04 log ₁₀
<i>C. albicans</i> +100µl of sheep's blood	3	900 s	H ₂ O ₂	CH ₃ COOH	<8 log ₁₀

<i>B.subtilis</i> spores +100µl of sheep's blood	3	480 s	H ₂ O ₂	-	3 log ₁₀
<i>B.subtilis</i> spores +100µl of sheep's blood	5	600 s	H ₂ O ₂	CH ₃ COOH	≥3 log ₁₀
<i>B.subtilis</i> spores +100µl of sheep's blood	4	600 s	H ₂ O ₂	CH ₃ COOH	≥6 log ₁₀
<i>B.subtilis</i> spores +100µl of sheep's blood	4	900 s	H ₂ O ₂	CH ₃ COOH	≥6 log ₁₀
<i>B.subtilis</i> spores +100µl of sheep's blood	3	900 s	H ₂ O ₂	CH ₃ COOH	4 log ₁₀
<i>B.stearoth</i> spores +100µl of sheep's blood	5	480 s	H ₂ O ₂	-	<4 log ₁₀
<i>B.stearoth</i> spores +100µl of sheep's blood	4	600 s	H ₂ O ₂	CH ₃ COOH	≥5 log ₁₀
<i>B.stearoth</i> spores +100µl of sheep's blood	4	600 s	H ₂ O ₂	CH ₃ COOH	≥5 log ₁₀
<i>B.stearoth</i> spores +100µl of sheep's blood	3	900 s	H ₂ O ₂	CH ₃ COOH	≥5 log ₁₀

The table shows that the germicide potency of the method according to the invention is markedly improved in comparison with the method where

5 only the germicide compound H₂O₂ is used in combination with O₃.

Claims

1. A method for disinfecting and sterilizing microbial contaminated or infectious materials, e.g. medical instruments, waste from medical laboratories or hospitals, the steps comprising:
 - 5 (a) loading the material for disinfecting and sterilizing into a vacuum-proof sterilization chamber
 - (b) introducing liquid hydrogen peroxide into the sterilization chamber so as to penetrate the material;
 - (c) introducing liquid acetic acid into the sterilization chamber so as to penetrate
10 the material;
 - (d) evacuating gas from the sterilization chamber so as to evaporate the liquid at least partially;
 - (e) introducing gaseous ozone into the sterilization chamber;
 - (f) treating the material in a sterilization chamber for a sufficient time period so
15 as to disinfect and sterilize the materials
2. The method of claim 1 wherein the steps (b) and (c) are carried out in any sequence under *in situ* generation of peracetic acid.
3. The method of claim 1 wherein the steps (b) and (c) are carried out simultaneously under *in situ* generation of peracetic acid.
- 20 4. The method according to one of the claims 1 to 3 wherein the method is carried out at a temperature of about 15 °C to 35 °C.
5. The method of claim 4 wherein the method is carried out at ambient temperature.
6. The method of claim 2 or 4 wherein the hydrogen peroxide and
25 the acetic acid of steps (b) and (c) are introduced in two separate ampoules having such properties that after the evacuation of step (d) they burst with release of their content.
7. The method of claim 3 or 4 wherein the hydrogen peroxide and the acetic acid of steps (b) and (c) are introduced in a container having at least
30 two compartments being constructed in such a manner that their contents are released is during the evacuation of step (d).

8. The method of claim 7 wherein the container has a compartment with a catalyst for shifting the equilibrium of the reaction equation
$$\text{H}_2\text{O}_2 + \text{CH}_3\text{-C(O)OH} \rightleftharpoons \text{H}_2\text{O} + \text{CH}_3\text{-C(O)OOH}$$
to the right, which catalyst is released at the same time as the hydrogen
5 peroxide and the acetic acid.
9. The method of claim 8 wherein the catalyst is sulfuric acid.
10. The method according to one of the claims 1 to 9 wherein in step (f) the ozone and the formed peracetic acid are present in the vapor phase.
11. The method according to one of the claims 1 to 10 wherein the
10 steps (b) to (e) are carried out at least two times.
12. The method according to one of the claims 1 to 11 wherein at the sterilization chamber feeding means for feeding hydrogen peroxide, acetic acid and ozone are disposed.
13. The method according to one of the claims 1 to 12 wherein the
15 molar ratio of acetic acid to ozone is 3/1 to 1/3.
14. The method according to one of the claims 1 to 13 wherein the humidity in step (e) or (f) is at least 10 %.
15. The method according to one of the claims 1 to 14 wherein the material is infectious waste in a solid or liquid state.
- 20 16. The method according to one of the claims 2 to 15 for treating contaminated or infectious waste wherein the waste is ground before, during or after step (b) or (c).
17. The method according to one of the claims 1 to 16 wherein step (e) is carried out at least twice.

INTERNATIONAL SEARCH REPORT

International Application No

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A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 A61L2/20 A61L11/00 //A61L101/10,A61L101/36

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 A61L

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	WO 94 21120 A (ENVIRO MEDICAL SYSTEMS INC) 29 September 1994 (1994-09-29) page 8, line 21 - line 29 page 9, line 10 - line 17 page 9, line 27 -page 10, line 4 page 11, line 23 - line 27 ---	1-5, 7-12, 14-17
A	US 5 700 426 A (BARDAT ANNIE ET AL) 23 December 1997 (1997-12-23) column 2, line 35 - line 60 column 3, line 1 - line 3 claims 1,4,5 --- -/--	1-5, 7-12, 14-17



Further documents are listed in the continuation of box C.



Patent family members are listed in annex.

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C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	<p>US 5 567 444 A (HEI ROBERT D ET AL) 22 October 1996 (1996-10-22) column 1, line 54 -column 2, line 20 column 5, line 30 - line 59 column 10, line 10 - line 35 claims 1-7</p>	1-17
A	<p>US 5 674 450 A (SWANZY JAMES ARCHIE ET AL) 7 October 1997 (1997-10-07) column 4, line 38 - line 52 column 5, line 9 - line 36</p>	1-17

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Information on patent family members

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